



Europäisches Patentamt
European Patent Office
Office européen des brevets

⑪ Publication number:

0 395 083
A2

⑫

EUROPEAN PATENT APPLICATION

⑬ Application number: **90108049.9**

⑮ Int. Cl.⁵: **C08F 4/02, C08F 4/654**

⑯ Date of filing: **27.04.90**

⑭ Priority: **28.04.89 IT 2032989**

⑮ Date of publication of application:
31.10.90 Bulletin 90/44

⑯ Designated Contracting States:
AT BE CH DE ES FR GB IT LI NL SE

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⑳ Components and catalysts for the polymerization of olefins.

㉑ Spherical catalyst components for the polymerization of olefins comprising a titanium compound and optionally an electron-donor supported on anhydrous magnesium chloride, characterized by a surface area between 20 and 250 m²/g, a porosity between 0.25 and 0.5 cc/g, and an X-ray spectrum where the magnesium chloride reflections are present at 2 ϑ of 35° and 14.95°, or where the reflection at 35° is substituted by a halo with the maximum intensity between 33.5° and 35° and the reflection of 2 ϑ 14.95° is absent.

EP 0 395 083 A2

Components And Catalysts For The Polymerization Of Olefins

This invention refers to catalyst components for the polymerization of olefins, the catalysts obtained from them and the use of same in the polymerization of olefins such as ethylene, propylene and their mixtures.

5 Catalysts for the polymerization of olefins comprising a titanium halide supported on anhydrous magnesium halides in active form are described extensively in patent literature.

Patent literature has become abundant since the use of magnesium halides in active form as supports for Ziegler-Natta catalyst components was described for the first time in USA patents n. 4,298,718 and 4,495,338.

10 The most active forms of the magnesium halides are characterized by X-ray spectra where the maximum intensity reflection which appears in the spectrum of the nonactive halides is no longer present, but is substituted by a halo with the maximum intensity shifted towards the lower angles with respect to the one of the maximum intensity reflection in the nonactive halide.

15 In the less active forms of the magnesium chloride the maximum intensity reflection which appears at 2.56 Å (2 = ϑ 35°) is no longer present, but is substituted by a halo with a maximum intensity between the angles 2 ϑ of 33.5° and 35°; a reflection at 2 ϑ of 14.95° is always present.

20 The introduction in industrial practice of catalysts supported on magnesium chloride made possible significant simplifications to the polyolefin production processes. In particular, the possibility of obtaining catalysts in the form of spherical particles capable of producing polymers which duplicate the form of the catalyst, have satisfactory morphological characteristics (flowability and bulk density), and do not require granulation which, as is well known, is expensive in terms of energy use.

Examples of catalysts with controlled particle size are described in USP 3,953,414.

25 The polymer (polyethylene) which can be obtained with said catalysts has good morphological characteristics: the polymer productivity of these catalysts, however, is not very high (generally between 2,000 and 15,000 g/g catalyst). When the polymer yield rises to values higher than 20,000 g/g catalyst, the polymer particles formed are fragile and the apparent density is very low.

30 The catalyst components described in the above mentioned US patent are obtained from an MgCl₂.6H₂O adduct spherulized in an apparatus of the dry cooling type, and then reacted with TiCl₄.

35 USP 4,399,054 describes catalyst components for the polymerization of olefins capable of producing polymer (polypropylene) with good flowability and bulk density characteristics. The polymer productivity of the catalyst is not very high (between 3,000 and 9,000 g/g catalyst; polymerization in heptane at 70 °C for 4 h with partial propylene pressure of 7 atm.).

40 The catalyst components are obtained from MgCl₂ adducts with alcohols, in the form of spherical particles containing generally 3 moles of alcohol.

45 Before the reaction with TiCl₄ the alcohol content is lowered to 2.5-2 moles in order to render the catalysts suitable to obtain non-fragile spherical form polymers. The alcohol content is never lowered below the 2 moles (this reduces drastically the activity of the catalyst).

50 In the case of magnesium chloride, at least in the less active forms (those where in the spectrum are present two halos with intensity peaks respectively between 2 ϑ angles from 30.45° to 31°, and from 33.5° to 35°), the reflection that in the nonactive magnesium chloride spectrum appears at 2 ϑ of 14.95° is still present.

55 Catalyst components, for the polymerization of CH₂ = CHR olefins, where R is hydrogen or an alkyl or aryl radical with 1-8 carbon atoms have now been found, suitable to obtain catalysts capable of producing polymers in the form of spherical particles with optimum morphological characteristics (flowability and high bulk density). Moreover, the catalysts have a significant catalytic activity and stereospecificity.

60 The catalyst components of the invention are in the form of spherical particles having an average diameter between 10 and 350 μ m, and comprise a titanium compound containing at least a Ti-halogen link, and optionally an electron-donor compound, supported on magnesium chloride. They are characterized by a surface area between 20 and 250 m²/g, porosity greater than 0.2 cc/g and comprised between 0.2 and 0.5 cc/g, X-ray spectrum (CuK α) where

65 a) reflections at 2 ϑ of 35° and 14.95° (characteristic of magnesium chloride) are present, or
b) where the reflection at 2 ϑ of 35° is substituted by a halo with a maximum intensity between the 2 ϑ angles of 33.5° and 35°, and the reflection at 2 ϑ of 14.95° is not present.

70 For the definition of the X-ray spectrum of magnesium chloride, reference is made to the standard ASTM D-3854. The registration of the spectrum is done using a copper anti-cathode and K α radiation.

75 The spectrum with the a) characteristic is peculiar of the catalytic components with surface area smaller

than 70-80 m²/g and porosity greater than 0.4 cc/g. The spectrum with the b) characteristic is given by components with an area greater than 60 m²/g and porosity between 0.25 and 0.4 cc/g.

The pore volume distribution is such that more than 50% of the pores have a radius greater than 100 Å. In the components with area smaller than 100 m²/g, more than 70% of the pores have a radius greater than 100 Å.

As already indicated, the catalyst components of the invention supply catalysts suitable for the production of olefins (co)polymers in the form of spherical particles with valued morphologic characteristics (high bulk density values, flowability, and mechanical resistance). The average diameter of the polymeric particles is between 50 and 5000 µm.

In particular, catalysts obtained from components with surface area smaller than 100 m²/g and a porosity greater than 0.4 cc/g, are appropriately used in the preparation of ethylene polymers (HDPE and LLDPE). The catalysts have a very high activity and the spherical polymer obtained has attractive morphological characteristics (very high bulk density, flowability and mechanical resistance).

The catalysts obtained from components with surface area greater than 60-70 m²/g and porosity smaller than 0.4 cc/g are preferred for use in preparing crystalline propylene homo and copolymers, the so called impact copolymers obtained with sequential polymerization of 1) propylene, and 2) ethylene-propylene mixtures.

They are also advantageously used in the preparation of ethylene-propylene rubbers (EP rubbers), or ethylene-propylene-diene rubbers (EPDM rubbers), and of propylene polymer compositions which contain said rubbers.

It is surprising that with the catalysts of this invention said types of rubber in spherical particles with good flowability and bulk density characteristics can be obtained, because up to now it has not been possible to prepare elastomeric polymers of the above indicated type in flowable granular particles due to the insurmountable problems of the fouling of the reactors and/or the agglomeration of the particles.

Particularly in the case of polypropylene, using stereospecific catalysts obtained from components with a surface area around 60-70 m²/g, porosity lower than 0.4 cc/g, and an X-ray spectrum of the type b), it is possible to obtain crystalline propylene homopolymers and propylene-ethylene copolymers, containing lesser portions of ethylene, characterized by very elevated porosity values which render them very attractive for the preparation of masterbatch with pigments and/or additives.

It is also surprising that the catalysts of the invention are highly active although the magnesium chloride in them gives X-ray spectra characteristic of the low active forms of the magnesium chloride itself.

Finally it is surprising and completely unexpected that the magnesium chloride is present in the crystalline form with an X-ray spectrum as the one indicated in b).

The preparation of the catalyst components is carried out in a variety of ways. The preferred method consists in starting with magnesium chloride/alcohol adducts containing alcohol moles in such a number that the adduct is solid at room temperature, but melted at temperatures between 100-130 °C.

The number of alcohol moles vary with the different types of alcohol.

The alcohols suitable for use have the ROH formula where R is an alkyl, cycloalkyl, or aryl radical with 1-12 carbon atoms. It is also possible to use mixtures of said alcohols.

Examples of alcohols are methanol, ethanol, propanol, butanol, 2-ethylhexanol and their mixtures.

With alcohols such as ethanol, propanol, and butanol, the number of moles is about 3 per MgCl₂ moles. The alcohol and the magnesium chloride are mixed in an inert hydrocarbon liquid immiscible with the adduct, brought to the fusion temperature of the adduct. The mixture undergoes vigorous stirring [using for example an ULTRA TURRAX T-45N apparatus rotating at 2000-5000 RPM (Jonke & Kunkel K.G.IKG Werke)].

The emulsion obtained is cooled in a very short time. This causes the solidification of the adduct in the form of spherical particles with the desired dimensions. The particles are dried and then subjected to a partial dealcoholation by heating it to temperatures going from 50 ° to 130 °C.

The partially dealcoholated adduct is in the form of spherical particles with an average diameter between 50 and 350 µm, a surface area between 10 and 50 m²/g and porosity from 0.6 to 2 cc/g (determined with a mercury porosimeter).

The higher the degree of dealcoholation, the higher the porosity. The pore volume distribution is such that more than 50% of the pores have a radius greater than 10,000 Å.

The dealcoholation is carried out until the alcohol contents are not greater than 2 moles per MgCl₂ mole, preferably comprised between 0.15 and 1.5 moles, particularly between 0.3 and 1.5 moles.

Whenever the dealcoholation is brought to values lower than 0.2 alcohol moles per MgCl₂ mole, the catalytic activity is considerably reduced.

The partially dealcoholated adduct is then suspended in cold TiCl₄ at a concentration of 40-50 g/l, and

then brought to a temperature of 80°-135°C and maintained at said temperature for 0.5-2 hrs. The excess $TiCl_4$ is separated hot via filtration or sedimentation.

The treatment with $TiCl_4$ is repeated one or more times if the alcohol content desired should be very low (generally less than 0.5% by weight).

5 During the preparation of a catalytic component containing an electron-donor compound, the latter is added to the $TiCl_4$ in quantities equal to molar ratios with respect to $MgCl_2$ between 1:6 and 1:16.

After the treatment with $TiCl_4$, the solid is washed with a hydrocarbon (e.g. hexane or heptane) and then dried.

10 According to another method, the melted adduct while in emulsion in an inert hydrocarbon is passed through a tube of the proper length under turbulent motion and then gathered in an inert hydrocarbon maintained at low temperature. Said method is described in USP n. 4,399,054 to which we make reference for the description. In this case as well the particles of the adduct are subjected to partial dealcoholization and reaction with $TiCl_4$.

15 As a variant to the methods described above the titanium compound, especially when it is solid at room temperature such as for instance the $TiCl_3$, is dissolved in the melted adduct which is then dealcoholated as indicated above and subjected to a reaction with a halogenating agent capable of reacting and separating the hydroxyl groups, such as for instance $SiCl_4$.

In the starting melted adduct, besides the titanium compound and optionally other transition metals, also co-supports such as $AlCl_3$, $AlBr_3$, $ZnCl_2$ may be included.

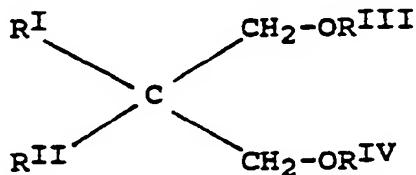
20 The titanium compounds suitable for the preparation of catalyst components, besides $TiCl_4$ and $TiCl_3$ and similar halides, also include other compounds with at least one Ti-halogen link, such as halogen alcoholates like trichlorophenoxytitanium and trichlorobutoxytitanium.

Finally the titanium compound can be used in mixtures with other transition metal compounds, such as V, Zr and Hf halides and halogen alcoholates.

25 As already indicated, the catalyst component may also contain an electron-donor compound (internal donor). This is necessary when the catalyst component must be used in the stereoregular polymerization of olefins such as propylene, 1-butene and 4-methyl-1-pentene.

The electron-donor compounds can be selected among compounds including ethers, esters, amines, ketons.

30 Preferred compounds are the alkyl esters, cycloalkyls and aryls of polycarboxyl acids, such as phthalic and maleic acids, and ethers of the formula



40 where R^1 , R^2 , same or different from each other, are alkyl, cycloalkyl or aryl radicals with 1-18 carbon atoms; R^3 and R^4 , same or different, are alkyl radicals with 1-4 carbon atoms.

Ethers of this type are described in U.S. patent application Serial No. 359,234, filed May 31, 1989 corresponding to EP-A-344 755 (European patent application No. 89 109 856), the disclosure of which is 45 incorporated herein by reference.

Examples representing said compounds are n-butyl phthalate, diisobutyl phthalate, di-n-octyl phthalate, 2-methyl-2-isopropyl-1,3-dimethoxypropane; 2-methyl-2-isobutyl-1,3-dimethoxypropane, 2,2-diisobutyl-1,3-dimethoxypropane, 2-isopropyl-2-isopentyl-1,3-dimethoxypropane.

The internal donor is generally present in molar ratios with respect to Mg of 1:8-1:14. The titanium 50 compound expressed as Ti is present in a percentage between 0.5 and 10% by weight.

Al-alkyl compounds selected particularly among Al-trialkyls, such as Al-triethyl, Al-triisobutyl and Al-tri-n-butyl, are used as co-catalysts.

The Al/Ti ratio is greater than 1 and is generally between 20-800.

In case of stereoregular polymerization of alpha olefins such as propylene and 1-butene, besides the Al-alkyl compound, usually, also an electron-donor compound (external donor is used). This compound may be the same as or different from the electron-donor compound present as internal donor.

When the internal donor is an ester of a polycarboxylic acid, especially a phthalate, the external donor is preferably selected among the silicon compounds of the formula $R_1 R_2 Si(OR)_2$, where R_1 and R_2 are

alkyl, cycloalkyl or aryl radicals with 1-18 carbon atoms, and R is an alkyl radical with 1-4 carbon atoms. Examples representing these silanes are methyl cyclohexyl dimethoxysilane, diphenyl dimethoxysilane, methyl-t.butyl dimethoxysilane.

1,3-diethers with the formula previously shown may also be used advantageously.

5 If the internal donor is one of these diethers, there is no need to use an external donor, since the stereospecificity of the catalyst is in itself sufficiently high.

Catalysts containing an internal donor are used in the preparation of LLDPE with restricted molecular weight distribution. As already indicated, the catalysts are used in the polymerization of $\text{CH}_2 = \text{CHR}$ olefins where R is a hydrogen, or an alkyl or aryl radical with 1-8 carbon atoms, and mixtures of said olefins with or 10 without a diene.

The polymerization is done according to known methods operating in liquid phase, in the presence or absence of an inert hydrocarbon diluent, or in gaseous phase.

It is also possible to use liquid-gas mixed processes where in one or more stages the polymerization is carried out in liquid phase, and in one or more successive stages it is carried out in gaseous phase.

15 The polymerization temperatures are in general between 20° and 150° C, preferably between 60° and 90° C. The operation takes place at atmospheric pressure or higher.

The data shown in examples and text relative to the following properties have been determined according to the methods indicated below.

	<u>Property</u>	<u>Method</u>
20	- MIL flow index	ASTM-D 1238
25	- MIE " "	ASTM-D 1238
	- MIF " "	ASTM-D 1238
30	- Fraction soluble in xylene	(see Determination before Examples)
35	- Isotacticity Index (I.I.)	Weight percentage of polymer insoluble in xylene at 25°C.
40		Basically it conforms

45

50

55

5 with the weight
percentage of polymer
insoluble in boiling n-
heptane)

10 - Surface area B.E.T. (apparatus used
SORPTOMATIC 1800 - C.
15 Erba).

20 - Porosity Unless otherwise
indicated, the porosity
is determined with the
25 B.E.T. method (see
above). It is
calculated from the
30 integral pore
distribution curve in
function of the pores
35 themselves.

40 - Porosity (mercury) Determined by immersing
a known quantity of the
sample in a known
45 quantity of mercury
inside a dilatometer

50

55

and gradually
5 hydraulically
increasing the pressure
of the mercury. The
10 pressure of
introduction of the
mercury in the pores is
15 in function of the
diameter of same. The
20 measurement is done
using a porosimeter
25 "Porosimeter 2000
Series" (C. Erba). The
30 total porosity is
calculated from the
volume decrease of the
35 mercury and the values
of the pressure
40 applied.
- Bulk density DIN-53194
- Flowability
45 The time taken by 100 g
of polymer to flow
50
55

5

10

15

- Morphology

through a funnel with an output hole of a diameter of 1.25 cm, and walls with an inclination of 20° on the vertical.

ASTM-D 1921-63

Determination of percentage in soluble xylene

20

2 g of polymer are dissolved in 250 ml of xylene at 135 °C while agitating. After 20 minutes the solution is left to cool, still under agitation, until it reaches 25 °C.

After 30 minutes the precipitated material is filtered through filter paper; the solution is evaporated in nitrogen current and the residue is dried under vacuum at 80 °C until it reaches constant weight.

25

Thus the percentage of polymer soluble in xylene at room temperature is calculated.

EXAMPLES

30

PREPARATION OF MgCl₂/ALCOHOL ADDUCTS.

The MgCl₂/alcohol adducts in spherical particle form are prepared following the method described in example 2 of USP n. 4,399,054, but operating at 3,000 RPM instead of 10,000 RPM.

35

The adduct was partially dealcoholated by heat at increasing temperatures from 30 °C to 180 °C operating in nitrogen current.

PREPARATION OF THE SOLID CATALYST COMPONENT.

40

In a 1 litre flask equipped with a condenser and mechanical agitator were introduced, under a nitrogen current, 625 ml of TiCl₄. At 0 °C while agitating were added 25 g of partially dealcoholated adduct. It was then heated up to 100 °C in 1 hour, when the temperature reaches 40 °C diisobutylphthalate (DIBF) was added in molar ratio Mg/DIBF = 8.

45

The temperature was maintained at 100 °C for 2 hours, it was then left to decant and afterwards the hot liquid was syphoned off. 550 ml of TiCl₄ were added and it was heated to 120 °C for 1 hour. Finally it was left to settle and the liquid was syphoned off while hot; the residual solid was washed 6 times with 200 ml aliquote of anhydrous hexane at 60 °C and 3 times at room temperature.

50

POLYMERIZATION OF PROPYLENE

55

In a 4 l stainless steel autoclave equipped with an agitator and a thermostatic system, which had been degassed with nitrogen at 70 °C for 1 hour and then with propylene, was introduced at 30 °C without agitation but under light propylene flow, the catalytic system consisting of a suspension of the above solid catalyst component in 80 ml of hexane, 0.76 g of Al-triethyl and 8.1 mg of diphenyldimethoxysilane (DPMS). The suspension was prepared immediately prior to the test.

The autoclave was then closed and 1 NI of H_2 was introduced. While agitating 1.2 kg of liquid propylene were charged and the temperature was brought to 70 °C in five minutes, maintaining the value constant for 2 hours.

At the end of the test, the agitation was stopped and any unreacted propylene was removed. Once the 5 autoclave was cooled to room temperature, the polymer was recovered and then dried at 70 °C under nitrogen current in oven for 3 hours and then analyzed.

COPOLYMERIZATION OF ETHYLENE WITH BUTENE-1 (LLDPE)

10 The autoclave described above was degassed with propane instead of propylene. The catalytic system consisting of 25 cc of hexane, 1.05 g of Al-triisobutyl and the above catalyst component, was introduced in the autoclave at room temperature under light propane flow. The pressure was increased 5.5 atm with H_2 and then 2 atm with ethylene, prepolymerizing the ethylene until 15 g of ethylene were used (45 °C).

15 The propane and the hydrogen were degassed and after washing with H_2 the gaseous phase was formed with 37.0 g of ethylene, 31.9 g of butene-1, and 1.8 atm of H_2 (total pressure 15 atm).

An ethylene-butene-1 mixture was then fed in a weight ratio of 9:1 at 70 °C for 2 hours

At the end the autoclave was degassed and cooled rapidly to room temperature.

The recovered copolymer was dried at 70 °C in nitrogen for 4 hours in an oven.

20

POLYMERIZATION OF ETHYLENE

25 A 2.5 l stainless steel autoclave equipped with agitator and thermostatic system was flushed as described above for the test with propylene but using ethylene instead of propylene.

At 45 °C were introduced in H_2 current, 900 ml of a solution containing 0.5 g/l of Al-triisobutyl in anhydrous hexane and immediately afterwards, the catalyst component suspended in 100 ml of the above mentioned solution.

30 The temperature was rapidly brought to 70 °C and H_2 was fed until the pressure reached 3 atm., then ethylene up to 10.5 atm. These conditions were maintained for 3 hours replenishing continuously the ethylene depleted. At the end of the polymerization reaction, the autoclave was rapidly vented and cooled at room temperature.

The polymeric suspension was filtered and the solid residue dried in nitrogen at 60 °C for 8 hours.

35

Example 1

A spherical $MgCl_2 \cdot 3EtOH$ adduct (obtained as indicated in the general method) was dealcoholated until an $EtOH/MgCl_2$ molar ratio of 1.7 was obtained.

40 A product with the following characteristics was obtained:

- porosity (mercury) = 0.904 cc/g;
- surface area = 9.2 m²/g;
- bulk density = 0.607 g/cc.

45 From this adduct, through the $TiCl_4$ treatment described in the general method, a solid catalyst component in spherical form was obtained with the following characteristics:

- Ti = 2.5% by weight;
- DIBF = 8.2% by weight;
- porosity = 0.405 cc/g;
- surface area = 249 m²/g;
- bulk density = 0.554 g/cc.

50 The X-ray spectrum of this component did not have reflections at 2 ϑ of 14.95 °; a halo with a maximum intensity of 2 ϑ of 34.72 ° was present instead.

This catalyst component was used in the polymerization of propylene following the technique described in the general section. Using 0.01 g of component 430 g of polymer were obtained with the following 55 characteristics:

- fraction soluble in xylene at 25 °C = 2.4%;
- MIL = 2.5 g/10';
- bulk density = 0.48 g/cc;

- morphology: 100% spherical particles with diameter between 1000 and 5000 μm ;
- flowability: 10 sec.

5 Example 2

By partially dealcoholating (according to example 1) a $\text{MgCl}_2 \cdot 3\text{EtOH}$ spherical adduct also obtained according to the method indicated in example 1, an adduct was prepared with an $\text{EtOH}/\text{MgCl}_2$ molar ratio of 1.5, and with the following characteristics:

- 10 - porosity (mercury) = 0.946 cc/g;
- surface area = 9.1 m^2/g ;
- bulk density = 0.564 g/cc.

From this adduct, through TiCl_4 treatment as indicated above, a spherical catalyst component was prepared with the following characteristics:

- 15 - Ti = 2.5% weight;
- DIBF = 8.0% weight;
- porosity = 0.389 cc/g;
- surface area = 221 m^2/g ;
- bulk density = 0.555 g/cc.

20 The X-ray spectrum of the component did not show reflections at 2ϑ of 14.95° ; only a halo was present with a maximum intensity at 2ϑ of 2.5780° .

This catalyst component was used in the polymerization of the propylene using the procedure of example 1.

Using 0.015 g of catalyst component, 378 g of polypropylene were obtained with the following 25 characteristics:

- fraction soluble in xylene at 25°C = 2.6%;
- MIL - 2.8 g/10';
- bulk density = 0.395 g/cc;
- morphology = 100% spherical particles with diameter between 1000 and 5000 μm ;
- 30 - flowability - 12 sec.

Example 3

35 By partially dealcoholating (according to example 1) a $\text{MgCl}_2 \cdot 3\text{EtOH}$ spherical adduct obtained according to the method indicated in the preceding examples, an $\text{EtOH}/\text{MgCl}_2 = 1$ adduct was obtained which has the following characteristics:

- porosity (mercury) = 1.208 cc/g;
- surface area = 11.5 m^2/g ;
- 40 - bulk density = 0.535 g/cc.

From said adduct, through TiCl_4 reaction, following the methodology described in the preceding examples, was obtained a spherical catalytic component with the following characteristics:

- Ti = 2.2% weight;
- DIBF = 6.8% weight;
- 45 - porosity = 0.261 cc/g;
- surface area = 66.5 m^2/g ;
- apparent density = 0.440 g/cc.

The X-ray spectrum of the catalytic component showed a reflection at 2ϑ of 14.95° as well as one at $2\vartheta = 35^\circ$.

50 Using 0.023 g of the catalyst component in the propylene polymerization using the conditions in example 1, 412 g of polypropylene were obtained with the following characteristics:

- fraction soluble in xylene at room temperature = 3.0%;
- MIL - 3.2 g/10';
- bulk density = 0.35 g/cc;
- 55 - morphology = 100% spherical particles with diameter between 500 and 5000 μm ;
- flowability - 12 sec.

Following the previous general description of the procedure for the copolymerization of ethylene with butene, 0.0238 g of catalyst component were used and 240 g of copolymer was obtained with the following

characteristics:

- linked butene = 8.3% weight;
- fraction soluble in xylene at room temperature = 12.2%;
- MIE = 12 g/10';
- 5 - MIF = 12 g/10';
- MIF/MIE = 30;
- morphology = 100% spherical particles with diameter between 500 and 5000 μm .

10 Example 4

By partially dealcoholating (according to example 1) a $\text{MgCl}_2.3\text{EtOH}$ spherical adduct obtained according to the method indicated in the preceding examples, an $\text{EtOH}/\text{Mg} = 0.4$ adduct was obtained which had the following characteristics:

- 15 - porosity (mercury) = 1.604 cc/g;
- surface area = 36.3 m^2/g ;
- apparent density = 0.410 g/cc.

By treating this support with TiCl_4 at a temperature of 135 °C, at a concentration of 50 g/l, with three 1 hour treatments, a spherical catalyst component was obtained which, after the elimination of the excess 20 TiCl_4 , washing and drying, had the following characteristics:

- $\text{Ti} = 2.6\%$ weight;
- porosity = 0.427 cc/g;
- surface area 66.5 m^2/g .

The X-ray spectrum of this component showed a reflection at 2 ϑ of 14.95° as well as at 2 ϑ of 35°.

25 Using 0.012 g of catalyst component in the polymerization of ethylene according to the method described in the general section, 400 g of polyethylene were obtained with the following characteristics:

- MIE = 0.144 g/10';
- MIF = 8.87 g/10';
- MIF/MIE = 61.6
- 30 - morphology = 100% spherical particles with diameter between 1000 and 5000 μm ;
- flowability = 12 sec.;
- apparent density = 0.38 g/cc.

35 Example 5

By partially dealcoholating (as per example 1) a $\text{MgCl}_2.3\text{EtOH}$ spherical adduct obtained according to the method indicated in the preceding examples, an adduct was obtained with $\text{EtOH}/\text{MgCl}_2$ molar ratio of 0.15 with the following characteristics:

- 40 - porosity (mercury) = 1.613 cc/g;
- surface area = 22.2 m^2/g .

The X-ray spectrum of this component showed a reflection at 2 ϑ of 14.95° as well as 2 ϑ of 35°.

Using 0.03 g of this component in the polymerization of ethylene, as described in example 4, 380 g of polyethylene were obtained with the following characteristics:

- 45 - MIE = 0.205 g/10';
- MIF = 16.42 g/10';
- MIF/MIE = 80.1;
- flowability = 12 sec.;
- bulk density = 0.40 g/cc.

50

Example 6

55 A $\text{MgCl}_2.1\text{EtOH}$ adduct was prepared according to the method in example 3, but using also a quantity of water diluted in the alcohol used for the preparation of the starting $\text{MgCl}_2.3\text{EtOH}$ equal to 2% in weight.

The adduct after dealcoholation contained 3% in weight of water. With said adduct, after treatment with TiCl_4 and DIBF as described in example 1, a spherical catalyst component was obtained with the following weight composition:

- Ti = 2.35%;
- DIBF = 6.9%.

Using 0.025 g of this component in the polymerization of propylene as per example 1, 410 g of polymer in spherical particle form were obtained which had the following characteristics:

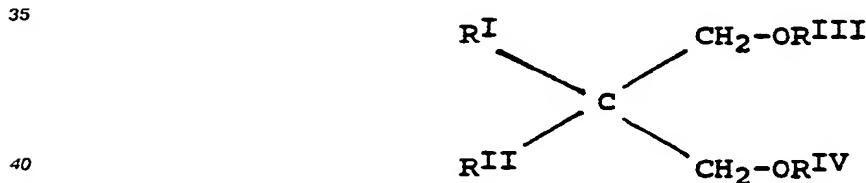
- 5 - fraction soluble in xylene at 25 °C = 3.1%;
- MIL = 3.0 10¹⁰;
- apparent density = 0.35 g/cc;
- morphology = 100% of the spherical particles having a diameter between 100 and 5000 µm ;
- flowability = 13 sec.

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Claims

1. A catalyst component for the polymerization of olefins comprising a titanium compound containing at least one Ti-Halogen link supported on anhydrous magnesium chloride, said component being in the form of spherical particles with an average diameter between 10 and 350 µm , surface area between 20 and 250 m²/g, porosity greater than 0.2 cc/g, having an X-ray spectrum where a) reflections at angle 2 ϑ of 35° and 2 ϑ of 14.95° are present, or b) where the reflection at 2 ϑ angle of 35° is no longer present but is substituted by a halo with a maximum intensity between angles 2 ϑ of 33.5° and 35°, and the reflection at angle 2 ϑ of 14.95° is not present.
2. The component of claim 1, further comprising an electron-donor compound in molar ratio with the magnesium chloride between 1:4 and 1:20.
3. The component of claim 1 or 2, where the surface area is less than 100 m²/g, the porosity is greater than 0.44 cc/g and having an X-ray spectrum of the type a).
- 25 4. The component of claim 1 or 2, where the area is greater than 60 m²/g, the porosity is between 0.2 and 0.4 cc/g, and having an X-ray spectrum of the type b).
5. The component of one or more of the preceding claims, where the pore volume distribution is such that at least 50% of them have a radius greater than 100 Å.
6. The component of one or more of the preceding claims, where the titanium compound is TiCl₄.
- 30 7. The component of one or more of claims 2 to 6, where the electron-donor compound is selected among alkyl, cycloalkyl, or aryl esters of phthalic acid.
8. The component of one or more of claims 2 to 6, where the electron-donor compound is selected among 1,3-diethers of the formula:

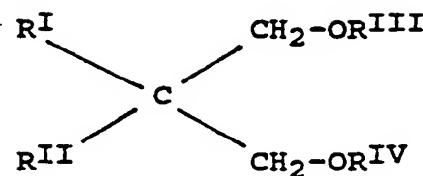
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where R¹, R², same or different from each other, are alkyl, cycloalkyl or aryl radicals with 1-18 carbon atoms, R³ and R⁴, same or different, are alkyl radicals with 1-4 carbon atoms.

- 45 9. The component of one or more of the preceding claims, where the titanium compound is present in quantity between 0.5% and 10% in weight, expressed as metallic Ti.
10. A catalyst for the polymerization of olefins, comprising the product of the reaction between a catalyst component of claim 1, and an Al-trialkyl compound.
- 50 11. The catalyst of claim 10, where an electron-donor compound (external donor) is used in the preparation of the catalyst.
12. The catalyst of claim 11, where the solid catalyst component includes an electron-donor selected among alkyl, cycloalkyl and aryl esters of phthalic acid, and the external donor is selected among silicon compounds of the formula R¹R²Si(OR)₂, where R¹ and R² equal or different are alkyl, cycloalkyl or aryl radicals with 1-18 carbon atoms, and R is an alkyl radical with 1-4 carbon atoms.
- 55 13. The catalyst of claim 11, where the external donor is selected among 1,3-diethers of the formula:



where R^I, R^{II}, same or different from each other, are alkyl, cycloalkyl or aryl radicals with 1-18 carbon atoms; and R^{III} and R^{IV}, same or different, are alkyl radicals with 1-4 carbon atoms.

10 14. A MgCl₂/ROH alcohol adduct, where R is an alkyl, cycloalkyl or alkylaryl radical with 1-12 carbon atoms, suitable for the preparation of the catalyst component of one or more of claims 1 to 9, containing from 0.2 to 2 moles of alcohol per mole of MgCl₂, and having a surface area between 10 and 50 m²/g, porosity (mercury) from 0.6 to 2.5 cc/g and pore volume distribution such that at least 50% of the pores 15 have a radius greater than 10,000 Å.

15 16. A polymer or copolymer of olefins in the form of spherical particles with an average diameter between 50 and 5000 µm, and obtained with catalysts described in claims 10 to 13.

16. Process for preparing the catalyst component of one or more of the preceding claims 1 to 9, comprising the reaction of:

20 1) a magnesium chloride/alcohol adduct containing alcohol moles in such a number that the adduct is solid at room temperature, but melted at temperatures between 100 and 130 °C and 2) with a titanium compound containing at least one halogen-titanium bond.

17. Process for the polymerization of olefins carried out in the presence of the catalyst of the preceding 25 claims 10 to 13.

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Europäisches Patentamt
European Patent Office
Office européen des brevets



⑪ Publication number:

0 395 083 A3

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EUROPEAN PATENT APPLICATION

⑬ Application number: **90108049.9**

⑮ Int. Cl.5: **C08F 4/02, C08F 4/654**

⑯ Date of filing: **27.04.90**

⑭ Priority: **28.04.89 IT 2032989**

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⑮ Date of publication of application:
31.10.90 Bulletin 90/44

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⑯ Designated Contracting States:
AT BE CH DE ES FR GB IT LI NL SE

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⑲ Components and catalysts for the polymerization of olefins.

⑳ Spherical catalyst components for the polymerization of olefins comprising a titanium compound and optionally an electron-donor supported on anhydrous magnesium chloride, characterized by a surface area between 20 and 250 m²/g, a porosity between 0.25 and 0.5 cc/g, and an X-ray spectrum where the magnesium chloride reflections are present at 2 θ of 35° and 14.95°, or where the reflection at 35° is substituted by a halo with the maximum intensity between 33.5° and 35° and the reflection of 2 θ 14.95° is absent.

EP 0 395 083 A3



EUROPEAN SEARCH
REPORT

Application Number

EP 90 10 8049

DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)						
Y	EP-A-0 065 700 (EUTECO IMPIANTI S.p.A.) * Claim 1 * - - -	1-17	C 08 F 4/02 C 08 F 4/654						
D,Y	US-A-4 399 054 (M. FERRARIS) * Example 2 * - - - - -	1-17							
The present search report has been drawn up for all claims									
<table border="1"><tr><td>Place of search</td><td>Date of completion of search</td><td>Examiner</td></tr><tr><td>The Hague</td><td>01 March 91</td><td>SERRAVALLE M.</td></tr></table>				Place of search	Date of completion of search	Examiner	The Hague	01 March 91	SERRAVALLE M.
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The Hague	01 March 91	SERRAVALLE M.							
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